

1	1011.6	97.5	168104	4	US-09-949-016-12026	Sequence	12026	<i>f</i>
2	1011.6	97.5	168105	4	US-09-949-016-16554	Sequence	16554	<i>f</i>
3	598.6	57.7	601	4	US-09-949-016-1703431	Sequence	170431	<i>f</i>
4	200	19.3	51620	4	US-09-949-016-120848	Sequence	12848	<i>f</i>
5	200	19.3	51621	4	US-09-949-016-165503	Sequence	16503	<i>f</i>
6	200	19.3	75799	4	US-09-949-016-15211	Sequence	15231	<i>f</i>
7	198.8	19.2	14345	4	US-09-949-016-15449	Sequence	15449	<i>f</i>
8	198.8	19.2	39243	4	US-09-949-016-12316	Sequence	12316	<i>f</i>
9	198.8	19.2	39243	4	US-09-949-016-15443	Sequence	15443	<i>f</i>
10	198.4	19.1	601	4	US-09-949-016-13148	Sequence	13148	<i>f</i>
11	198.2	19.1	88490	4	US-09-949-016-12758	Sequence	12758	<i>f</i>
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13	196.8	19.0	601	4	US-09-949-016-45906	Sequence	45906	<i>f</i>
14	196.8	19.0	20441	4	US-09-949-016-13052	Sequence	13052	<i>f</i>
15	196.4	18.9	601	4	US-09-949-016-15905	Sequence	45905	<i>f</i>
16	196.4	18.9	89689	4	US-09-949-016-13079	Sequence	13079	<i>f</i>
17	196.2	18.9	601	4	US-09-949-016-121170	Sequence	121170	<i>f</i>
18	196.2	18.9	601	4	US-09-949-016-127095	Sequence	127095	<i>f</i>
19	196.2	18.9	601	4	US-09-949-016-140139	Sequence	140139	<i>f</i>
20	196.2	18.9	71119	4	US-09-949-016-15538	Sequence	15358	<i>f</i>
21	196	18.9	73853	4	US-09-949-016-127029	Sequence	12029	<i>f</i>
22	195.8	18.9	601	4	US-09-949-016-127094	Sequence	127094	<i>f</i>
23	195.6	18.8	90618	4	US-09-949-016-15964	Sequence	15964	<i>f</i>
24	195.4	18.8	17132	4	US-09-949-016-15361	Sequence	15361	<i>f</i>
25	195.4	18.8	24150	4	US-09-949-016-12418	Sequence	12438	<i>f</i>
26	195.2	18.8	45427	4	US-09-949-016-16243	Sequence	16243	<i>f</i>
27	195	18.8	601	4	US-09-949-016-156268	Sequence	56268	<i>f</i>

C	28	1.5	18.8	1820.0	4	US-09-949-016-15660	A	Sequence	15660, A
C	29	1.95	18.8	1820.0	4	US-09-949-016-15661	A	Sequence	15661, A
C	30	1.95	18.8	1971.9	4	US-09-949-016-15662	A	Sequence	15662, A
C	31	1.95	18.8	1971.9	4	US-09-949-016-15663	A	Sequence	15663, A
C	32	1.95	18.8	5567.4	4	US-09-949-016-15263	A	Sequence	12563, A
C	33	1.95	18.8	5567.4	4	US-09-949-016-15106	A	Sequence	15766, A
C	34	1.95	18.8	7480.4	4	US-09-949-016-15118	A	Sequence	15118, A
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C	38	1.94.4	18.7	601	4	US-09-949-016-63026	A	Sequence	63026, A
C	39	1.94.4	18.7	601	4	US-09-949-016-63027	A	Sequence	63027, A
C	40	1.94.4	18.7	58545.3	4	US-09-949-016-13565	A	Sequence	13565, A
C	41	1.94.4	18.7	373318.2	4	US-09-949-016-17771	A	Sequence	17371, A
C	42	1.94.4	18.7	373368.4	4	US-09-949-016-12062	A	Sequence	12062, A
C	43	1.94.4	18.7	678583.3	4	US-09-949-016-14577	A	Sequence	14577, A
C	44	1.94.4	18.7	678583.3	4	US-09-949-016-14578	A	Sequence	14578, A
C	45	1.94	18.7	25401.4	4	US-09-949-016-13345	A	Sequence	13345, A

ALIGNMENTS

RESULT 1

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: Sequence 12026, Application US/05949016
: Patent No. 6812339
: GENERAL INFORMATION:
: APPLICANT: VENTER, J. Craig et al.
: TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
: FILE REFERENCE: CL001307
: CURRENT APPLICATION NUMBER: US/09/949,016
: PRIOR FILING DATE: 2000-04-14
: PRIOR APPLICATION NUMBER: 60/241,755
: PRIOR FILING DATE: 2000-10-20
: PRIOR APPLICATION NUMBER: 60/237,768
: PRIOR FILING DATE: 2000-10-03
: PRIOR APPLICATION NUMBER: 60/231,498
: PRIOR FILING DATE: 2000-09-08
: NUMBER OF SEQ ID NOS: 207012
: SOFTWARE: FastSeq for Windows Version 4.0
: SEQ ID NO 12026
: LENGTH: 168104
: TYPE: DNA
: ORGANISM: Human
: FEATURES:
: NAME/KEY: misc.feature
: LOCATION: (1)...(168104)
: OTHER INFORMATION: n = A,T,C or G
: US-09-949-016-12026

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Query Match	97.5%	Score 1011.6	DB 4	length 168104
Best Local Similarity	99.1%	Pred. No.8.5e-237		
Matches 1011, Conservative	5	Mismatches 4	Indels 0	Gaps 0

QY	16	TTAATGTAATAAATATTTGATATGAATTTAAAGGAAATTAGACATCATAGAAAAA	75
Db	82814	TTAATGTAATAAATATTTCTATATGAAATTTTAAGGAAATTAGACATCATAGAAAAA	82873
QY	76	TGCTCTTACTGTGAAACATTAATTTGTACATTTTGGTCAACTATCTTTCACTACTT	135
Db	82874	TGCTCTTACTGTGAAACATTAATTTGTACATTTTGGTCAACTATCTTTCACTACTT	82933
QY	136	TTTAGTACATTAATGTTAAGTTGTACAGTGGCAGTCTTATATGTAATATGGCAGCTGC	195
Db	82934	TTTAGTACATTAATGTTAAGTTGTACAGTGGCAGTCTTATATGTAATATGGCAGCTGC	82993
QY	196	AGCATGAAATATAACATATCTAATATTTTGTACTATCTTATAGGAAATATCAGAAATTT	255
Db	82994	AGCATGAAATATAACATATCTAATATTTTGTACTATCTTATAGGAAATATCAGAAATTT	83053
QY	256	CAAAACCTTGTAATTTTAAAGGTATAGTCACATTTTAAATATGTGCGGTATATTTATAC	315

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Qy      316 ATGATTTGACGTTTGTGMAAATATTTTCCCTGACCTTTATTTATAGATGATCTACAGT 375
Db      83114 ATGATTTGACGTTTGTGMAAATATTTTCCCTGACCTTTATTTATAGATGATCTACAGT 83173
Qy      376 GTAGGCAAACTTATATATGTCATCTCACTTATGTCATATGTCATATCTATCCCATG 435
Db      83174 GTAGGCAAACTTATATATGTCATCTCACTTATGTCATATGTCATATCTATCCCATG 83233
Qy      436 CTAAATTTATAGTTTGTAAAAATACGTTTGTAAATAGTTTGTAGTCTATATCA 495
Db      83234 CTAAATTTATAGTTTGTAAAAATACGTTTGTAAATAGTTTGTAGTCTATATCA 83293
Qy      496 AGCTTTCAAGGATTCATTTATAAAACTTTGKTTTATTTCTTGATATMCCGTTTT 555
Db      83294 AGCTTTCAAGGATTCATTTATAAAACTTTGKTTTATTTCTTGATATMCCGTTTT 83353
Qy      556 TCCATGCAAAAGTTAAATTTCTGACCTTTATTTTATTTATATATATAGATGAT 615
Db      83354 TCCATGCAAAAGTTAAATTTCTGACCTTTATTTTATATATATATAGATGAT 83413
Qy      616 GAGTATGACTACAAACAGGAAAAATTAACAGATTTGTTGTGCTTTGTCTAAATG 675
Db      83414 GAGTATGACTACAAACAGGAAAAATTAACAGATTTGTTGTGCTTTGTCTAAATG 83473
Qy      676 TTACTCTGACAAATCTTAAAGCAGTTCTTCACTTTGTTGATGATGATCTTACT 735
Db      83474 TTACTCTGACAAATCTTAAAGCAGTTCTTCACTTTGTTGATGATGATCTTACT 83533
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Db      83534 TAGTCCAGGGGCTGGGGCGATAGTGTGCTGTGCTCCAGTCTTTGCGGGCCGAG 83593
Qy      796 GCAGGTGATCACTTAAAGTCAGAGTTTGAACACAGCTGCCCAATGTTGAAAGTT 855
Db      83594 GCAGGTGATCACTTAAAGTCAGAGTTTGAACACAGCTGCCCAATGTTGAAAGTT 83653
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Db      83654 GTCTCTACTAAAAATACAAAAATTAAGACAGGCGTGTGGCACAATCTGTATTCAGCT 83713
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Db      83714 ACTCAGAGGCTTAACAAGGAAATCTTGAACCTGGGAGGAGGTTGCGATGAGCC 83773
Qy      976 ATTGCACTCAGGCTGGGCAACAGTGAAGTCTTGTCTCAAAAAAATTAATTAAT 1035
Db      83774 ATTGCACTCAGGCTGGGCAACAGTGAAGTCTTGTCTCAAAAAAATTAATTAAT 83833
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RESULT 2
US-09-949-016-16554
/ Sequence 16554, Application US/09949016
/ Patent No. 6812339
/ GENERAL INFORMATION:
/ APPLICANT: VENTER, J. Craig et al.
/ TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
/ WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
/ FILE REFERENCE: CL001307
/ CURRENT APPLICATION NUMBER: US/09/949,016
/ PRIOR FILING DATE: 2000-04-14
/ PRIOR APPLICATION NUMBER: 60/241,755
/ PRIOR FILING DATE: 2000-10-20
/ PRIOR APPLICATION NUMBER: 60/237,768
/ PRIOR FILING DATE: 2000-10-03
/ PRIOR APPLICATION NUMBER: 60/231,498
/ PRIOR FILING DATE: 2000-09-08
/ NUMBER OF SEQ ID NOS: 207012
/ SOFTWARE: FASTSEQ for Windows Version 4.0
/ SEQ ID NO 16554
/ LENGTH: 168105
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TYPE: DNA
ORGANISM: Human
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1) .. (168105)
OTHER INFORMATION: n = A, T, C or G
US-09-949-016-16554

Query Match 97.5%; Score 1011.6; DB 4; Length 168105;
Best Local Similarity 99.1%; Pred. No. 8.5e-237;
Matches 1011; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Qy      16 TTATGATTAATAATTTTCTATATGAAATTTTAAAGGAAATGAGCATCTAGAAAAA 75
Db      82814 TTATGATTAATAATTTTCTATATGAAATTTTAAAGGAAATGAGCATCTAGAAAAA 82873
Qy      76 TGCTTACTAGTGAACATTTATTTGTCATTTTGTGTACATTAATCTTCAATACTT 135
Db      82874 TGCTTACTAGTGAACATTTATTTGTCATTTTGTGTACATTAATCTTCAATACTT 82933
Qy      136 TTAGTAATTAATGTTAAGTTGATGACGAGTGTATATATATATAGTATGAGCTGAC 195
Db      82934 TTAGTAATTAATGTTAAGTTGATGACGAGTGTATATATATATAGTATGAGCTGAC 82993
Qy      196 AGCATGAAATTAACATATCTTAATTTTGTGACTATCTTATAGGAAATCAGAGAAATT 255
Db      82994 AGCATGAAATTAACATATCTTAATTTTGTGACTATCTTATAGGAAATCAGAGAAATT 83053
Qy      256 CAAAACCTTGTAGTTTATAGGATATAGTCAATTTTAAATGCGGTATATATAC 315
Db      83054 CAAAACCTTGTAGTTTATAGGATATAGTCAATTTTAAATGCGGTATATATATAC 83113
Qy      316 ATGATTTGACGTTTGTGMAAATATTTTCCCTGACCTTTATTTATAGATGATCTACAGT 375
Db      83114 ATGATTTGACGTTTGTGMAAATATTTTCCCTGACCTTTATTTATAGATGATCTACAGT 83173
Qy      376 GTAGGCAAACTTATATATGTCATCTCACTTATGTCATATGTCATATCTATCCCATG 435
Db      83174 GTAGGCAAACTTATATATGTCATCTCACTTATGTCATATGTCATATCTATCCCATG 83233
Qy      436 CTAAATTTATAGTTTGTAAAAATACGTTTGTAAATAGTTTGTAGTCTATATCA 495
Db      83234 CTAAATTTATAGTTTGTAAAAATACGTTTGTAAATAGTTTGTAGTCTATATCA 83293
Qy      496 AGCTTTCAAGGATTCATTTATAAAACTTTGKTTTATTTCTTGATATMCCGTTTT 555
Db      83294 AGCTTTCAAGGATTCATTTATAAAACTTTGKTTTATTTCTTGATATMCCGTTTT 83353
Qy      556 TCCATGCAAAAGTTAAATTTCTGACCTTTATTTTATTTATATATAGATGAT 615
Db      83354 TCCATGCAAAAGTTAAATTTCTGACCTTTATTTTATTTATATATAGATGAT 83413
Qy      616 GAGTATGACTACAAACAGGAAAAATTAACAGATTTGTTGTTGCTTTGCTAAATG 675
Db      83414 GAGTATGACTACAAACAGGAAAAATTAACAGATTTGTTGTTGCTTTGCTAAATG 83473
Qy      676 TTACTCTGACAAATCTTAAAGCAGTTCTTCACTTTGTTGATGATGATCTTACT 735
Db      83474 TTACTCTGACAAATCTTAAAGCAGTTCTTCACTTTGTTGATGATGATCTTACT 83533
Qy      736 TAGTCCAGGGGCTGGGGCGATAGTGTGCTGTGCTCCAGTCTTTGCGGGCCGAG 795
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Qy      796 GCAGGTGATCACTTAAAGTCAGAGTTTGAACACAGCTGCCCAATGTTGAAAGTT 855
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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 6, 2005, 06:38:25 ; Search time 121 Seconds

(Without alignments)
156.622 Million cell updates/sec

Title: US-10-664-358-549

Sequence: 1 MNLGLMFSCGLMLKLMKC.....ISFANSRSSEPTKMKMSFM 49

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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1: geneseqp1980s:*
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7: geneseqp2003bs:*
8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	260	100.0	49	3	AA87141 Human sec
2	260	100.0	49	4	AAE06118 Human gen
3	260	100.0	49	5	ABG33940 Human sec
4	260	100.0	49	6	ABR47655 Human sec
5	260	100.0	49	6	ABR00022 Human gen
6	260	100.0	49	7	ADB91468 Human sec
7	260	100.0	49	7	ADC74024 Human sec
8	260	100.0	50	2	AAW67873 Human sec
9	260	100.0	70	3	AA76058 Human skt
10	260	100.0	70	4	AA855997 Skin cell
11	260	100.0	70	4	AA806054 Human gen
12	260	100.0	70	5	ABG33876 Human sec
13	260	100.0	70	5	ABG33876 Human sec
14	260	100.0	71	3	AA872197 Human pro
15	260	100.0	106	2	AA59659 Secreted
16	260	100.0	106	2	AA59659 Secreted
17	260	100.0	106	3	AA59659 Secreted
18	260	100.0	106	5	AA804034 Human pro
19	260	100.0	106	5	AB889692 Human pol
20	260	100.0	106	7	ADW77809 Human pro
21	260	100.0	106	8	ADP19490 Human sec
22	260	100.0	112	3	AA843968 Human can
23	260	100.0	132	4	AB811978 Human can
24	217	83.5	108	4	AB85197 Drosophila
25	158	60.8	30	2	AA867999 Fragment

26	110	42.3	99	4	AAU32155 Novel hum
27	97	37.3	20	2	AAW67997 Fragment
28	73.5	28.3	107	3	AA809358 Arabidops
29	61	23.5	308	3	AA843188 Human ORF
30	61	23.5	360	7	ADB65455 Human pro
31	61	23.5	406	5	AD116911 NOX prot
32	61	23.5	415	5	AD116586 Human NOV
33	61	23.5	415	8	ADN42240 Human NOV
34	61	23.5	416	5	AD116588 Human NOV
35	61	23.5	416	8	ADN42242 Human NOV
36	59.5	22.9	57	3	AA809360 Arabidops
37	59.5	22.9	1970	4	AB864827 Drosophila
38	59	22.7	432	8	ADN18428 Bacterial
39	58.5	22.5	322	2	AAV30160 Human dor
40	58.5	22.5	733	4	ABG24605 Novel hum
41	58	22.3	234	4	ABG21806 Novel hum
42	58	22.3	391	7	ADB64644 Human pro
43	57.5	22.1	193	4	AA871944 Human Olf
44	57	21.9	358	5	ABP65847 Bifidobac
45	57	21.9	472	4	AB865583 Drosophila

ALIGNMENTS

RESULT 1
AA87141
ID AA87141 standard; protein; 49 AA.

AA87141;
DT 09-MAY-2000 (first entry)

Human secreted protein sequence SEQ ID NO:180.

Human; secreted protein; diagnosis; cytosolic; immunosuppressive;
antiinflammatory; nootropic; neuroprotective; anti-allergic; cancer;
tumour; neurodegenerative disorder; developmental abnormality; allergy;
foetal deficiency; blood disorder; immune system disorder; arthritis;
autoimmune disease; hepatic disease; renal disease; inflammation;
Alzheimer's disease; behavioural disorder; schizophrenia; osteoporosis;
infection; AIDS; spinal cord injury; transplant rejection; diabetes;
aetna; sepsis; acne; psoriasis; cardiovascular disorder;
reproductive disorder; gastrointestinal disorder; respiratory disorder;
metabolic disorder; food additive; preservative.

OS Homo sapiens.

XX WO200004140-A1.

XX 27-JAN-2000.

XX 14-JUL-1999; 99WO-US015649.

XX 15-JUL-1998; 98US-0092921P.

XX 15-JUL-1998; 98US-0092922P.

XX 15-JUL-1998; 98US-0092956P.

XX (HUMA-) HUMAN GENOME SCI INC.

XX Ruben SM, Komatsoulis G, Duan RD, Rosen CA, Moore PA, Shi Y,
PI Lafleur DW, Ehner R, Olsen HS, Brewer LA, Florence KA, Young PB;
PI Mucenaski M, Endress GA, Sopet DR,

DR WPI; 2000-161128/14.

XX N-PSDB; AA298094.

XX New isolated human genes, useful for diagnosis and treatment of, e.g.
PT cancers, neurological or blood disorders.

XX Claim 11; Page 429; 494pp; English.

XX The polynucleotide sequences given in AA298017 to AA298108 encode the

CC human secreted proteins given in AAY87064 to AAY87223. Human secreted
CC protein can have activities based on the tissues and cells the genes are
CC expressed in. Examples of activities include: cytostatic;
CC immunosuppressive; antiinflammatory; nocotropic; neuroprotective; and
CC antiallergic. The polynucleotides and their corresponding secreted
CC polypeptides are useful for preventing, treating or ameliorating medical
CC conditions, e.g. by protein or gene therapy. Also pathological conditions
CC can be diagnosed by determining the amount of the new polypeptides in a
CC sample or by determining the presence of mutations in the new
CC polynucleotides. Human secreted proteins and their polynucleotides can
CC be used for developing products for the diagnosis or treatment of cancer,
CC tumours, neurodegenerative disorders, developmental abnormalities and
CC foetal deficiencies, blood disorders, diseases of the immune system,
CC autoimmune diseases, hepatic and renal disease, inflammation, allergies,
CC Alzheimer's disease, behavioural disorders, schizophrenia, osteoporosis,
CC arthritis, infections, AIDS, spinal cord injuries, transplant rejection,
CC diabetes, asthma, sepsis, acne, psoriasis, cardiovascular disorders,
CC reproductive disorders, gastrointestinal disorders, respiratory disorders
CC and metabolic disorders. The proteins or polynucleotides can also be used
CC as food additives or preservatives. The proteins are also useful for
CC identifying their binding partners. AA298008 to AA298016 and AAY87063 are
CC sequence used in the exemplification of the present invention

XX Sequence 49 AA;

Query Match 100.0%; Score 260; DB 3; Length 49;
Best Local Similarity 100.0%; Pred. No. 2, 9e-30;
Matches 49; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNLGMIFSMCGMLKTKMCAMWAVYCSPIFANSRSSEPTKQMSFPM 49
DB 1 MNLGMIFSMCGMLKTKMCAMWAVYCSPIFANSRSSEPTKQMSFPM 49

RESULT 2

AAE06118 standard; protein; 49 AA.

AC AAE06118;

DT 24-SEP-2001 (first entry)

DE Human gene 14 encoded secreted protein HAUA183, SEQ ID NO:180.

KW Human; secreted protein; proliferative disorder; cancer; tumour; asthma;
KW foetal abnormality; developmental abnormality; haematopoietic disorder;
KW immune system disorder; AIDS; autoimmune disease; rheumatoid arthritis;
KW Parkinson's disease; cognitive disorder; schizophrenia; skin disorder;
KW psoriasis; sepsis; diabetes; atherosclerosis; cardiovascular disorder;
KW inflammation; neurological disorder; Alzheimer's disease; food additive;
KW angiotensin disorder; kidney disorder; gastrointestinal disorder; allergy;
KW pregnancy-related disorder; endocrine disorder; infection; wound healing;
KW cell culture; chemotaxis; vunerary; binding partner identification;
KW gene therapy; chromosome 19.

XX Homo sapiens.

OS Homo sapiens.

Key Location/Qualifiers
FT Peptide 1..37
FT /label= signal_peptide
FT Protein 38..49
FT /label= Mature_human_secreted_protein

MO200151504-A1.

19-JUL-2001.

12-JAN-2001; 2001MO-US000911.

PR 13-JAN-2000; 2000US-00482273.

XX (HUMA-) HUMAN GENOME SCI INC.

PI Ruben SM, Komatsuia GA, Duan DR, Rosen CA, Moore PA, Shi Y,
PI Lafleur DM, Olsen HS, Brewer LA, Florence KA, Young PE, Soppet DR,
PI Endress GA, Muscenski M, Ebner R;
XX WPI: 2001-425865/45.
DR N-PSDB; AAD11707.
XX
XX Isolated nucleic acid molecule encoding a human secreted protein is used
PT in preventing, treating or ameliorating a medical condition.

PS Claim 11; Page 796; 864pp; English.

XX AAD11330-AAD11721 represent cDNAs corresponding to 71 human secreted
CC protein genes, and AAB06041-AAB06132 represent the proteins they encode.
CC AAB06133-AAB06205 represent human secreted protein fragments. The
CC secreted proteins and their genes are useful for preventing, treating or
CC ameliorating medical conditions, e.g. by protein or gene therapy.
CC Pathological conditions can be diagnosed by determining the amount of the
CC new protein in a sample or by determining the presence of mutations in
CC the new genes. Specific uses are described for each of the 71 genes,
CC based on the tissues in which they are most highly expressed, and include
CC developing products for the diagnosis or treatment of proliferative
CC disorders, cancer, tumours, foetal and developmental abnormalities,
CC haematopoietic disorders, diseases of the immune system, AIDS, autoimmune
CC diseases (e.g., rheumatoid arthritis), inflammation, allergies,
CC neurological disorders (e.g., Alzheimer's disease, Parkinson's disease),
CC cognitive disorders, schizophrenia, asthma, skin disorders (e.g.,
CC psoriasis), sepsis, diabetes, atherosclerosis, cardiovascular disorders,
CC angiogenic disorders, kidney disorders, gastrointestinal disorders,
CC pregnancy-related disorders, endocrine disorders, and infections. The
CC proteins can also be used to aid wound healing and epithelial cell
CC proliferation, to prevent skin aging due to sunburn, to maintain organs
CC before transplantation, for supporting cell culture of primary tissues,
CC to regenerate tissues, to identify their cognate ligands or binding
CC partners, and in chemotaxis, and can be used as a food additive or
CC preservative to modify storage properties. Antibodies specific for a
CC protein of the invention can be used in alleviating symptoms associated
CC with the disorders mentioned above, and in diagnostic immunoassays (e.g.,
CC radioimmunoassay or enzyme linked immunosorbent assay (ELISA)). The
CC present sequence represents a human secreted protein of the invention

XX Sequence 49 AA;

Query Match 100.0%; Score 260; DB 4; Length 49;
Best Local Similarity 100.0%; Pred. No. 2, 9e-30;
Matches 49; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNLGMIFSMCGMLKTKMCAMWAVYCSPIFANSRSSEPTKQMSFPM 49
DB 1 MNLGMIFSMCGMLKTKMCAMWAVYCSPIFANSRSSEPTKQMSFPM 49

RESULT 3

ABG33940 standard; protein; 49 AA.

AC ABG33940;

DT 15-JUL-2002 (first entry)

DE Human secreted protein encoded by gene 14 #2.

KW Human; secreted protein; gene therapy; immunosuppressive; antiarthritic;
KW antirheumatic; antiproliferative; cytostatic; cardiant; vasotropic;
KW cerebroprotective; nocotropic; neuroprotective; antibacterial; vitricide;
KW fungicide; opthalmological; autoimmune disease; neoplasm;
KW rheumatoid arthritis; hyperproliferative disorder; cardiac arrest;
KW cardiovascular disorder; cerebrovascular disorder; cerebral ischemia;
KW angiotensin; nervous system disorder; Alzheimer's disease; infection;
KW ocular disorder; corneal infection; wound healing; skin aging;
KW epithelial cell proliferation; food additive.

XX Homo sapiens.